

Bulletin Number: MSA 20-81

**Distribution:** All Providers

**Issued:** February 2, 2021

**Subject:** COVID-19 Response: Coverage of U.S. Food & Drug Administration

(FDA) Emergency Use Authorization (EUA) Drugs, Devices and Biological Products for COVID-19 Prevention and Treatment

**Effective:** As Indicated

Programs Affected: Medicaid, Healthy Michigan Plan, MIChild, Maternity Outpatient Medical

Services (MOMS)

Note: Implementation of this policy is contingent upon approval of a State Plan Amendment (SPA) by the Centers for Medicare & Medicaid Services (CMS).

The purpose of this bulletin is to notify Michigan Department of Health and Human Services (MDHHS) Medicaid-enrolled providers of important information regarding the Medicaid program coverage and reimbursement of FDA EUA drugs, devices, and biological products for COVID-19 prevention and treatment.

### **COVID-19 EUA Drugs, Devices and Biological Products**

For the duration of the federally declared COVID-19 Public Health Emergency (PHE), Medicaid will cover FDA EUA drugs, devices, and biological products and their administration for COVID-19 prevention and treatment. The effective date and coverage parameters will be consistent with their respective FDA EUA status provisions.

During the PHE, when EUA drugs, devices, and biological products for COVID-19 prevention and treatment are procured and purchased by the federal government, they will be made available to Medicaid providers at no cost. Medicaid will not reimburse providers for drugs, devices, or products that are federally purchased or supplied for free. Providers may bill the procedure code and the cost of the EUA drug, device, or biological product as \$0.00.

If providers are required to purchase COVID-19 EUA drugs, devices and biological products at a future date, reimbursement will be in accordance with Medicaid policies specific to respective health care services described in the MDHHS Medicaid Provider Manual. The MDHHS Medicaid Provider Manual can be found on the MDHHS website at <a href="https://www.michigan.gov/medicaidproviders">www.michigan.gov/medicaidproviders</a> >> Policy, Letters and Forms.

## Administration of COVID-19 EUA Drugs, Devices and Biological Products

Coverage of provider administration services related to COVID-19 EUA drugs, devices and biological agents will be similar to existing administration services billed to Medicaid. Refer to the Practitioner Chapter (Injectable Drugs and Biological Products subsection) and the Billing & Reimbursement for Professionals Chapter (Ancillary Medical Services subsection) of the MDHHS Medicaid Provider Manual for information regarding billing and reimbursement for physician-administered drugs, devices and biological products. The MDHHS Medicaid Provider Manual can be found on the MDHHS website at <a href="https://www.michigan.gov/medicaidproviders">www.michigan.gov/medicaidproviders</a> >> Policy, Letters and Forms.

Reimbursement for the administration of COVID-19 EUA drugs, devices and biological products will be as follows:

- During the federally declared PHE period, including any extensions, reimbursement for the administration services for COVID-19 EUA drugs, devices and vaccine administration services will be temporarily increased to 100% of Medicare rates for equivalent services.
- Following the end of the federally declared PHE period, ongoing COVID-19 drugs, devices and biological product administration services will return to reimbursement rates consistent with existing Medicaid policies and State Plan methodology.

All reimbursement rates will be reviewed and updated as applicable and are published on the MDHHS website at <a href="www.michigan.gov/medicaidproviders">www.michigan.gov/medicaidproviders</a> >> Billing & Reimbursement >> Provider Specific Information. Additional pertinent coverage parameters are accessible via the Medicaid Code and Rate Reference tool within the Community Health Automated Medicaid Processing System (CHAMPS).

#### **Procedure Codes**

MDHHS will cover new Current Procedural Technology (CPT) codes for the reporting of EUA drugs, devices and biological products and their administration as they are announced during the COVID-19 PHE. Code coverage will have effective dates consistent with the FDA EUA dates. Providers are highly encouraged to review the latest CPT coding guidance on the new reporting structure and additional resources available at the Centers for Medicare & Medicaid Services (CMS) and American Medical Association (AMA).

# **Pharmacy Coverage**

For the duration of the COVID-19 PHE, EUA drugs approved for use for the treatment of COVID-19 in an outpatient setting will also be approved for pharmacy benefit coverage in accordance with existing policies and reimbursement rates that align with the practitioner fee screens as described above.

Any pharmacy-specific COVID-19 vaccine services billing instructions will be published at: <a href="https://michigan.magellanrx.com/provider/">https://michigan.magellanrx.com/provider/</a> and also incorporated into the Pharmacy Claims Processing Manual at: <a href="https://michigan.magellanrx.com/provider">https://michigan.magellanrx.com/provider</a> >> Documents >> Manuals.

#### **Additional Resources for Providers**

Information on FDA EUA drugs, devices and biological products for COVID-19 prevention and treatment can be found at: <a href="https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization">https://www.fda.gov/emergency-preparedness-and-response/mcm-legal-regulatory-and-policy-framework/emergency-use-authorization</a>.

#### **Public Comment**

The public comment portion of the policy promulgation process is being conducted concurrently with the implementation of the change noted in this bulletin. Any interested party wishing to comment on the change may do so by submitting comments to Rajita Dnyate, Policy Specialist, via e-mail at DnyateR@michigan.gov.

Please include "COVID-19 Response: Coverage of U.S. Food & Drug Administration (FDA) Emergency Use Authorization (EUA) Drugs, Devices and Biological Products for COVID-19 Prevention and Treatment" in the subject line.

Comments received will be considered for revisions to the change implemented by this bulletin.

#### **Manual Maintenance**

Information is time-limited and will not be incorporated into any policy or procedure manuals.

#### Questions

Any questions regarding this bulletin should be directed to Provider Inquiry, Department of Health and Human Services, P.O. Box 30731, Lansing, Michigan 48909-8231, or e-mailed to <a href="ProviderSupport@michigan.gov">ProviderSupport@michigan.gov</a>. When you submit questions, be sure to include your name, affiliation, NPI number, and phone number so you may be contacted if necessary. Providers may phone toll-free 1-800-292-2550.

**Approved** 

Kate Massey, Director

**Medical Services Administration**